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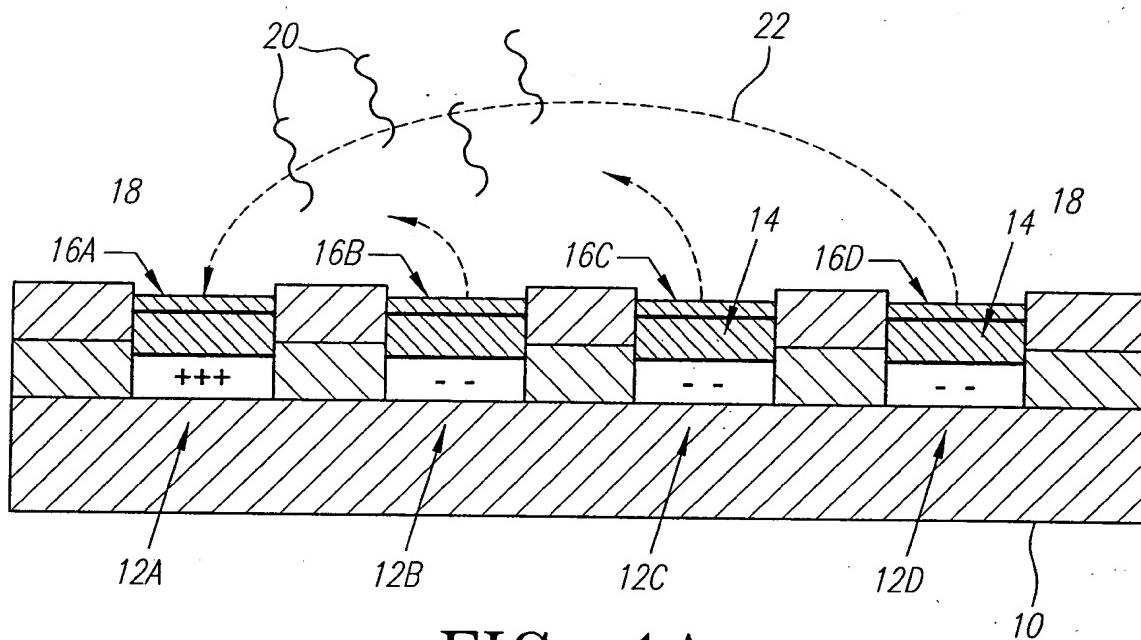


FIG. 1A

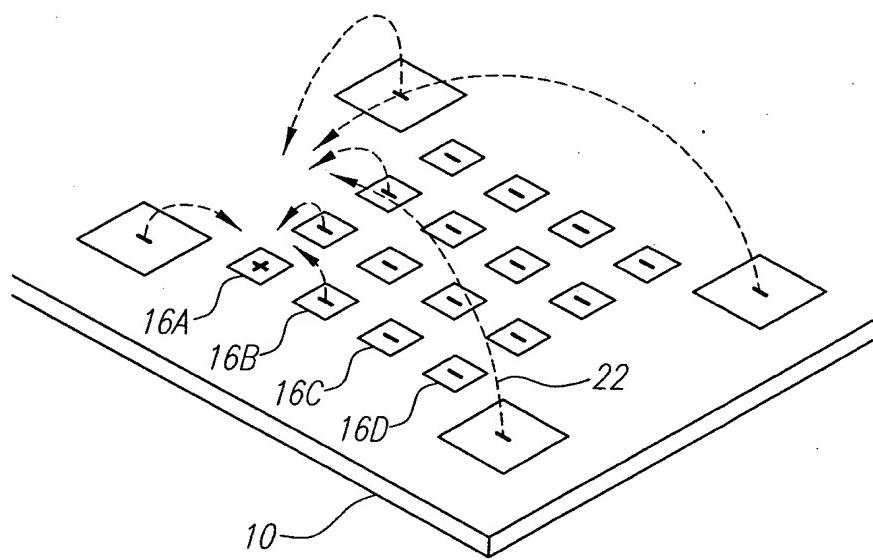
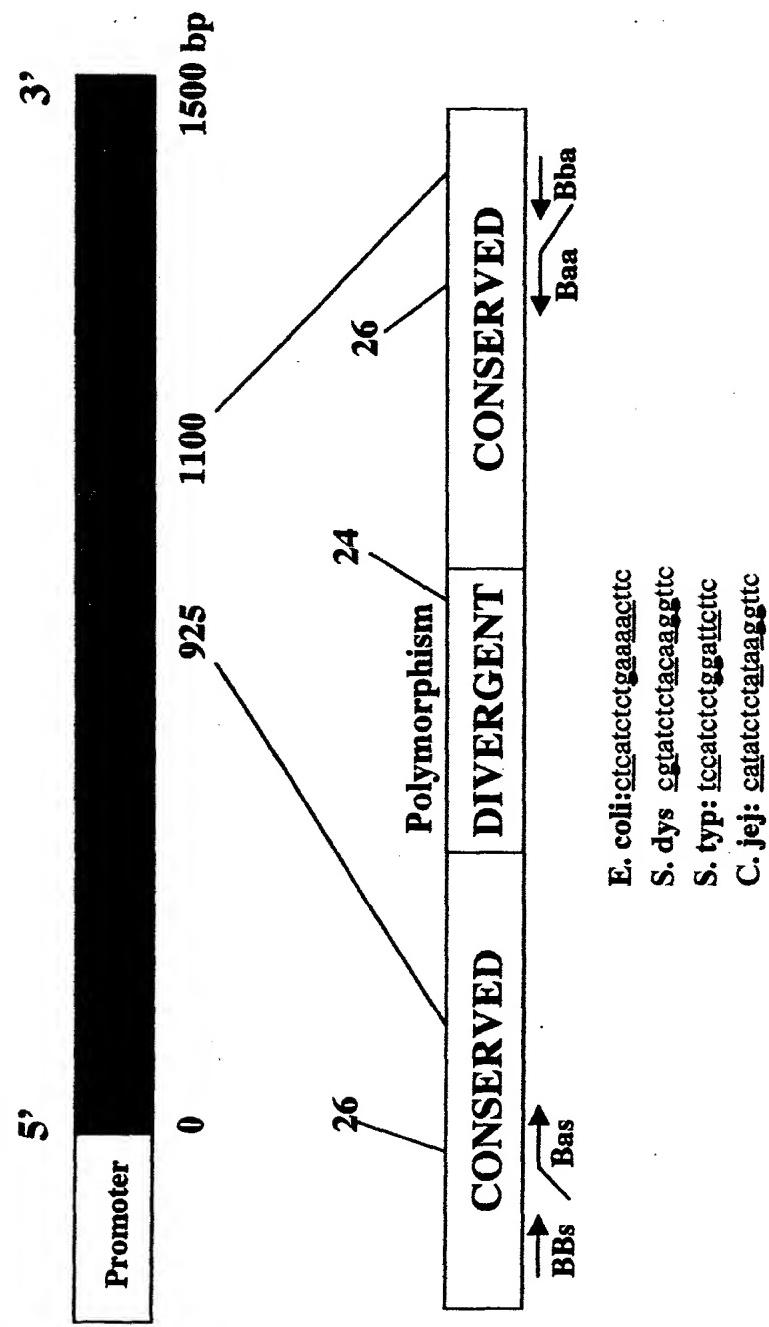


FIG. 1B

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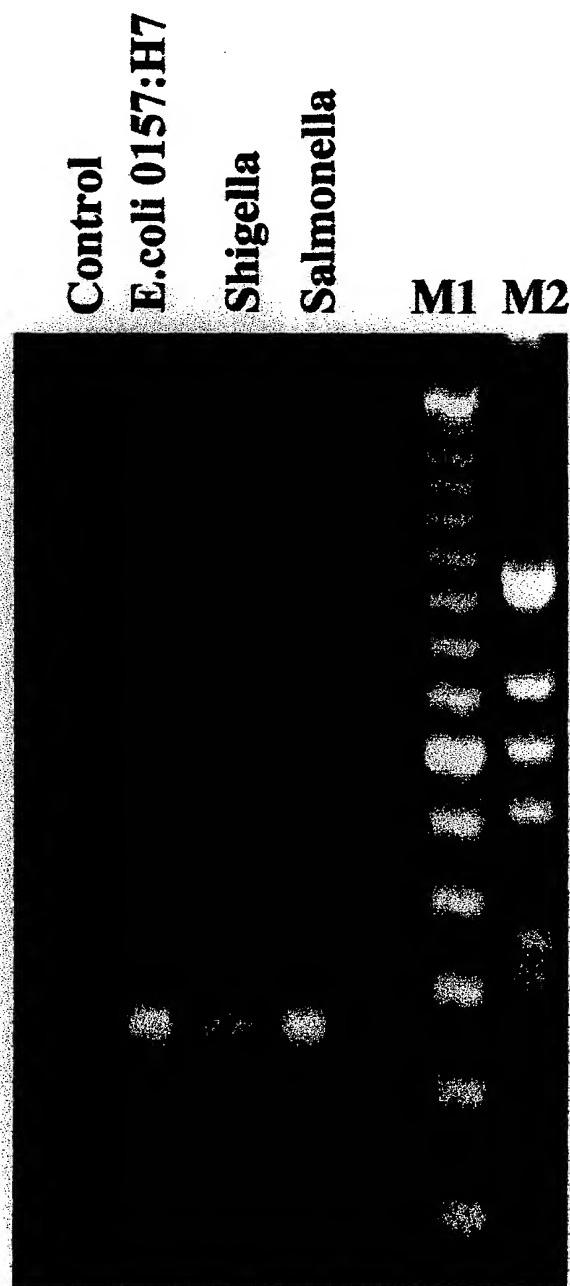
FIGURE 2A



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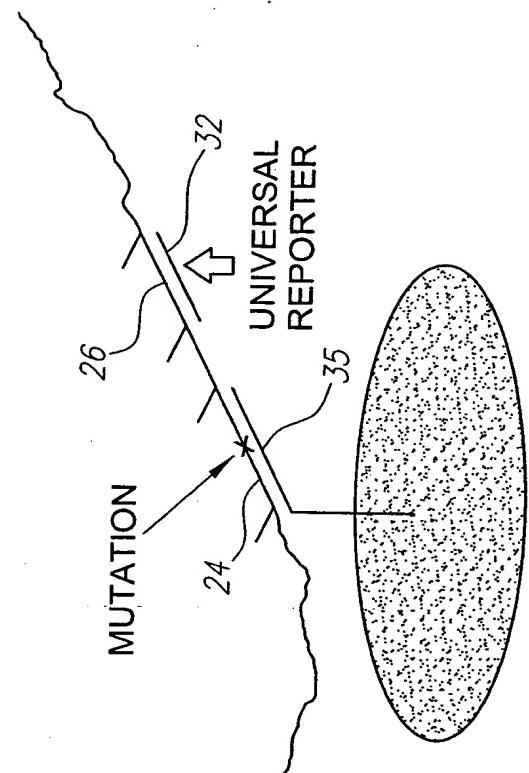
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FIGURE 2B

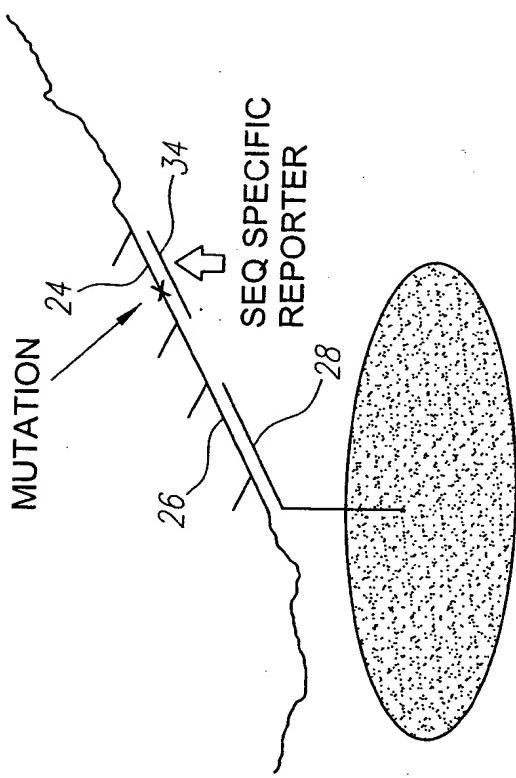


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SEQUENCE-SPECIFIC CAPTURE



UNIVERSAL CAPTURE

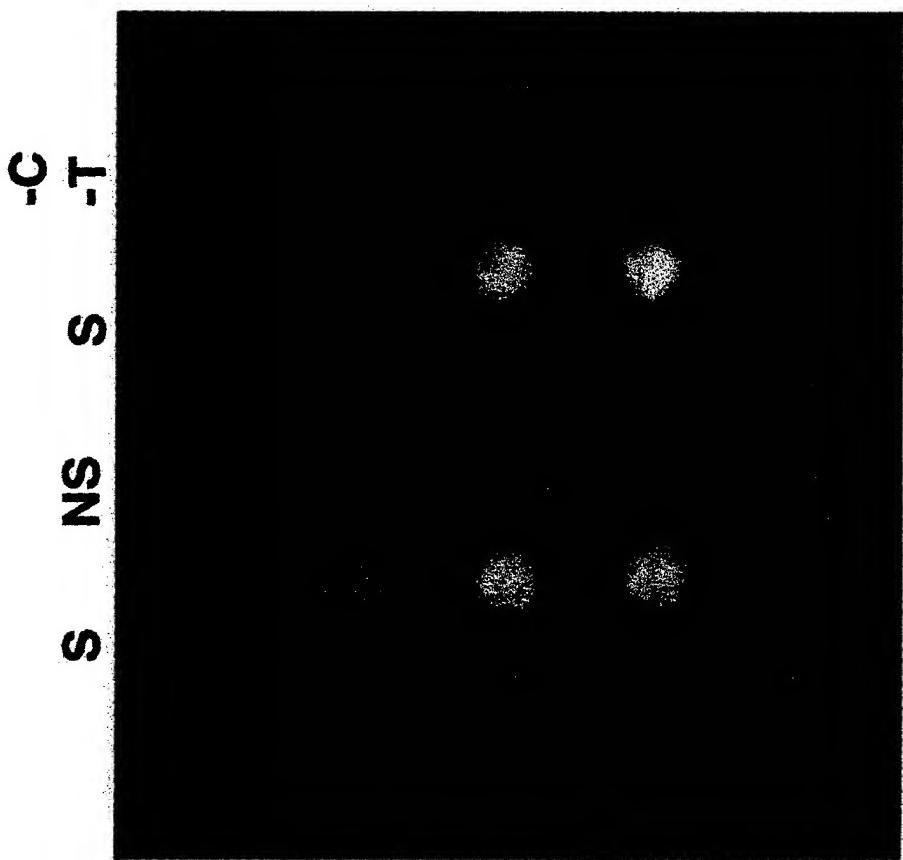
FIG. 2C

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FIG. 2D

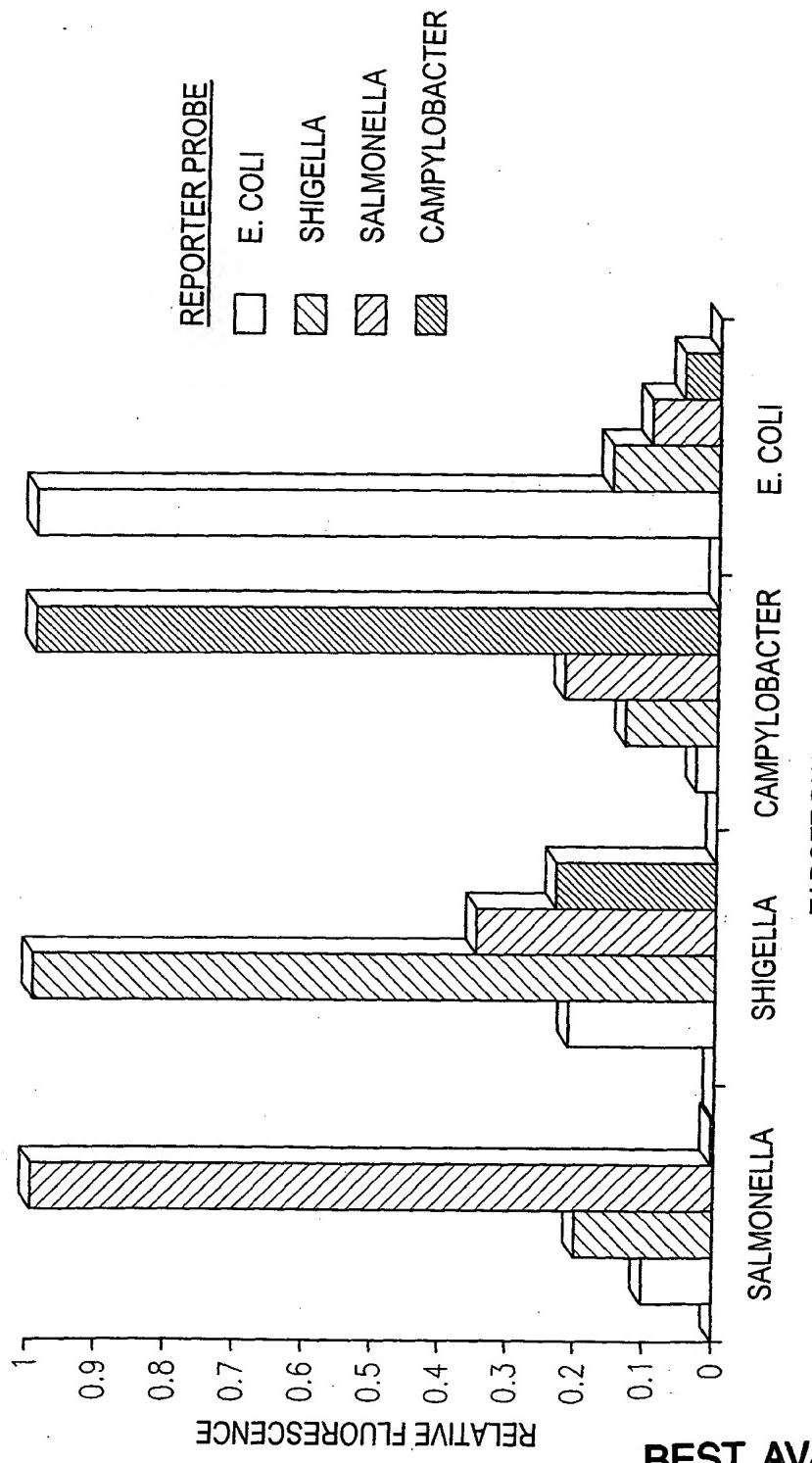
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FIGURE 3A



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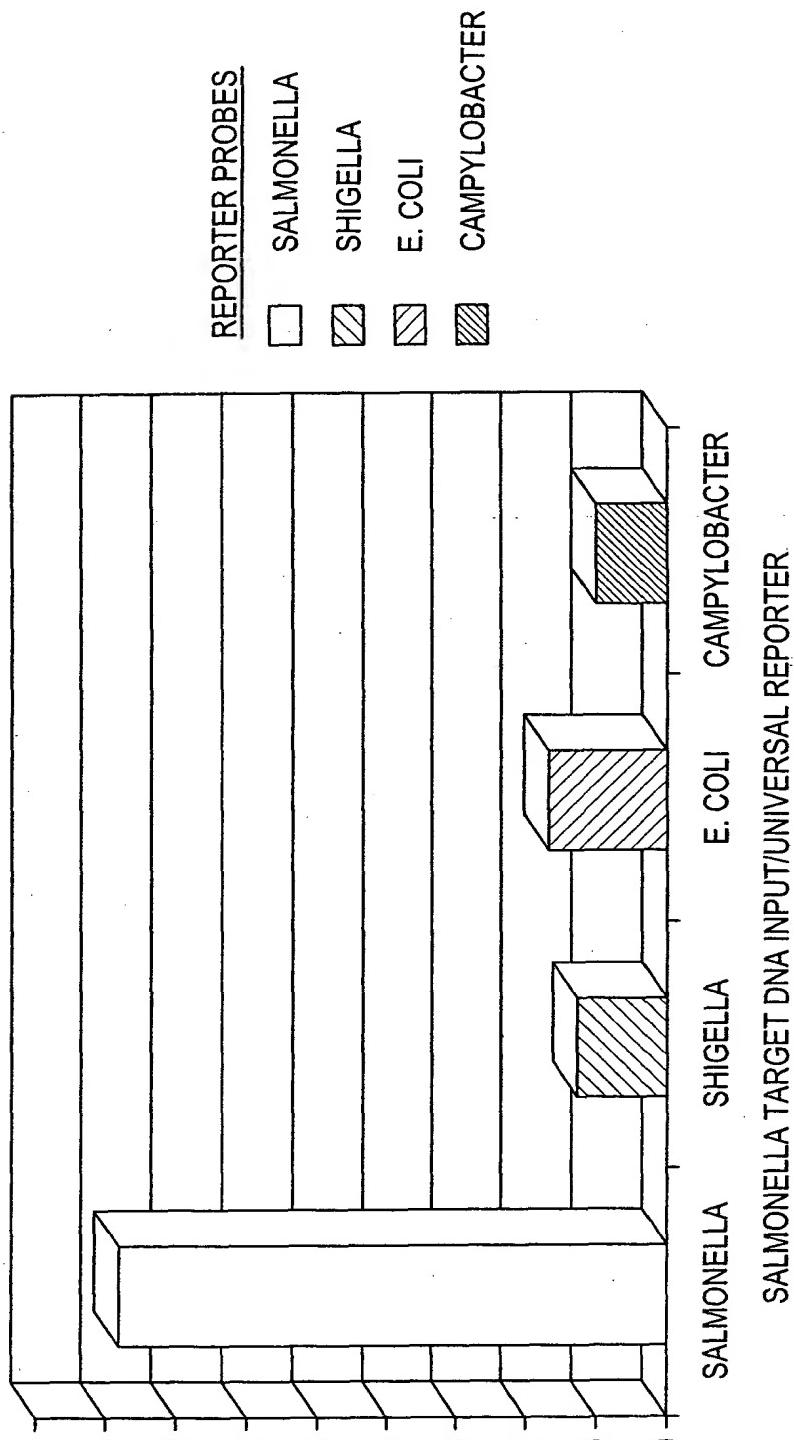
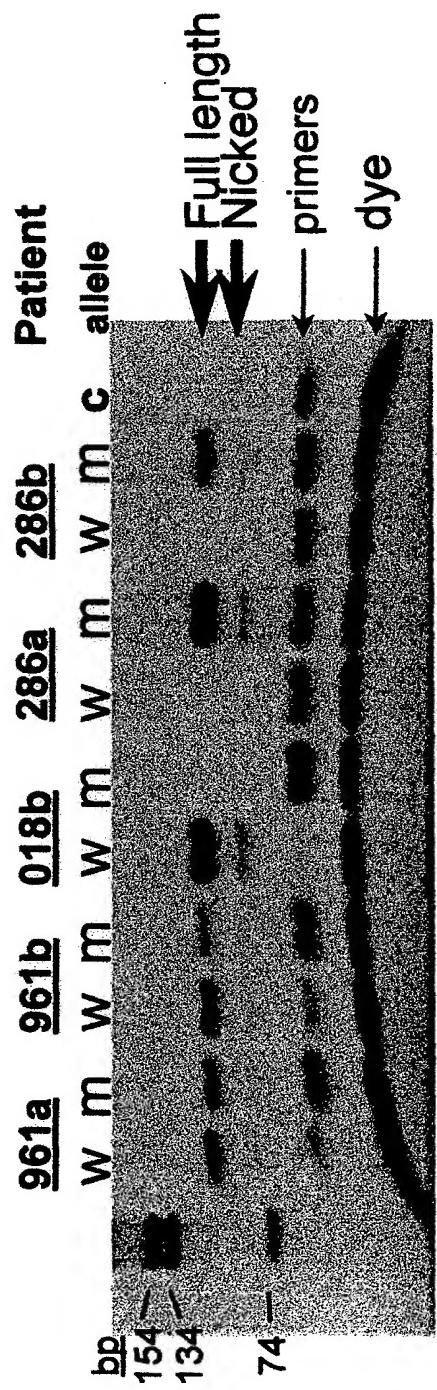


FIG. 3C

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FIGURE 4A



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W
M

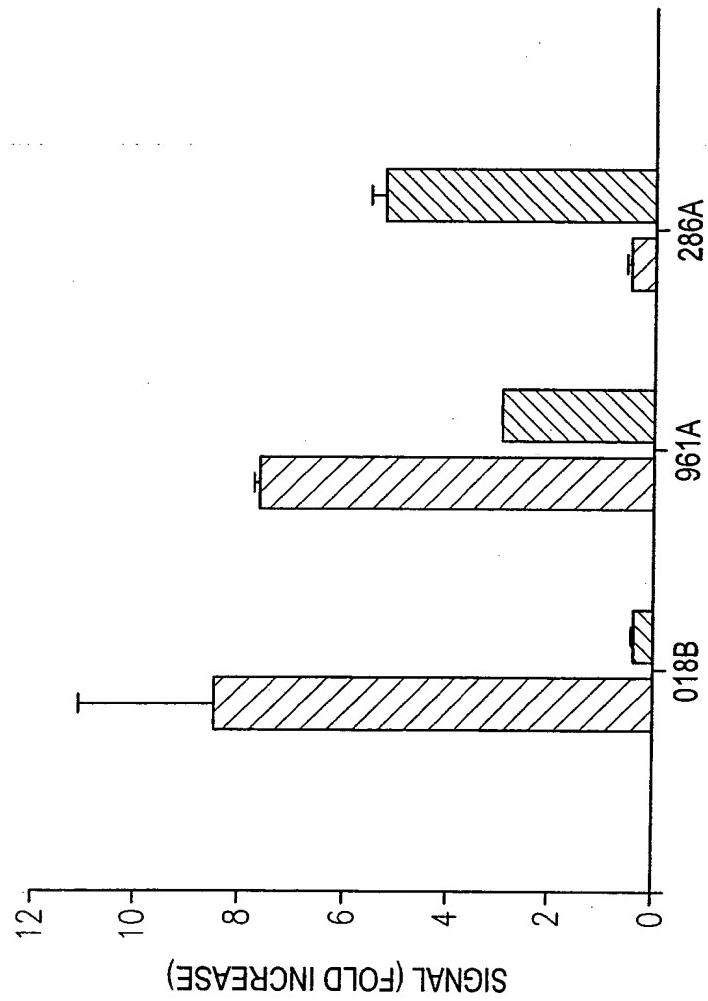


FIG. 4B

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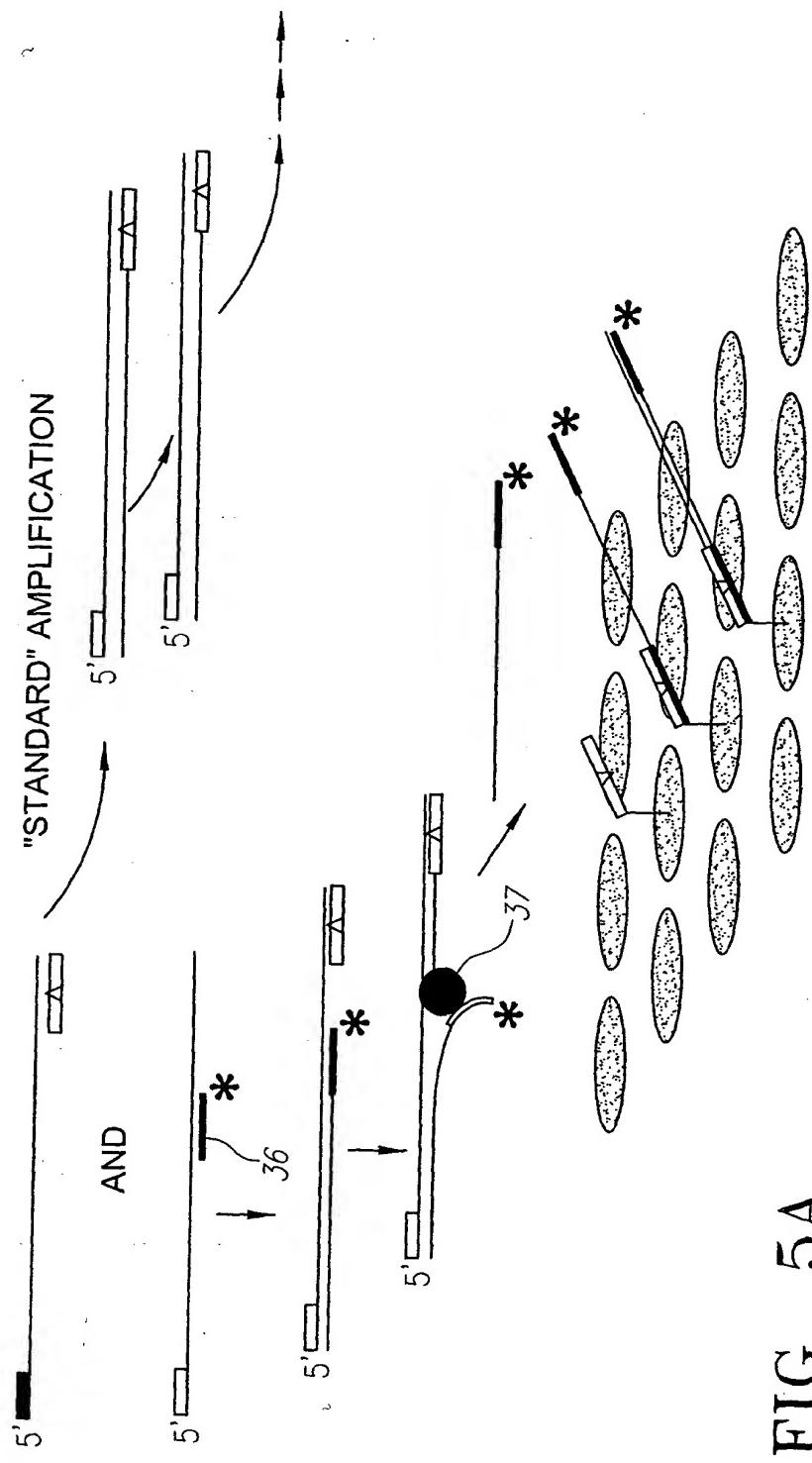


FIG. 5A

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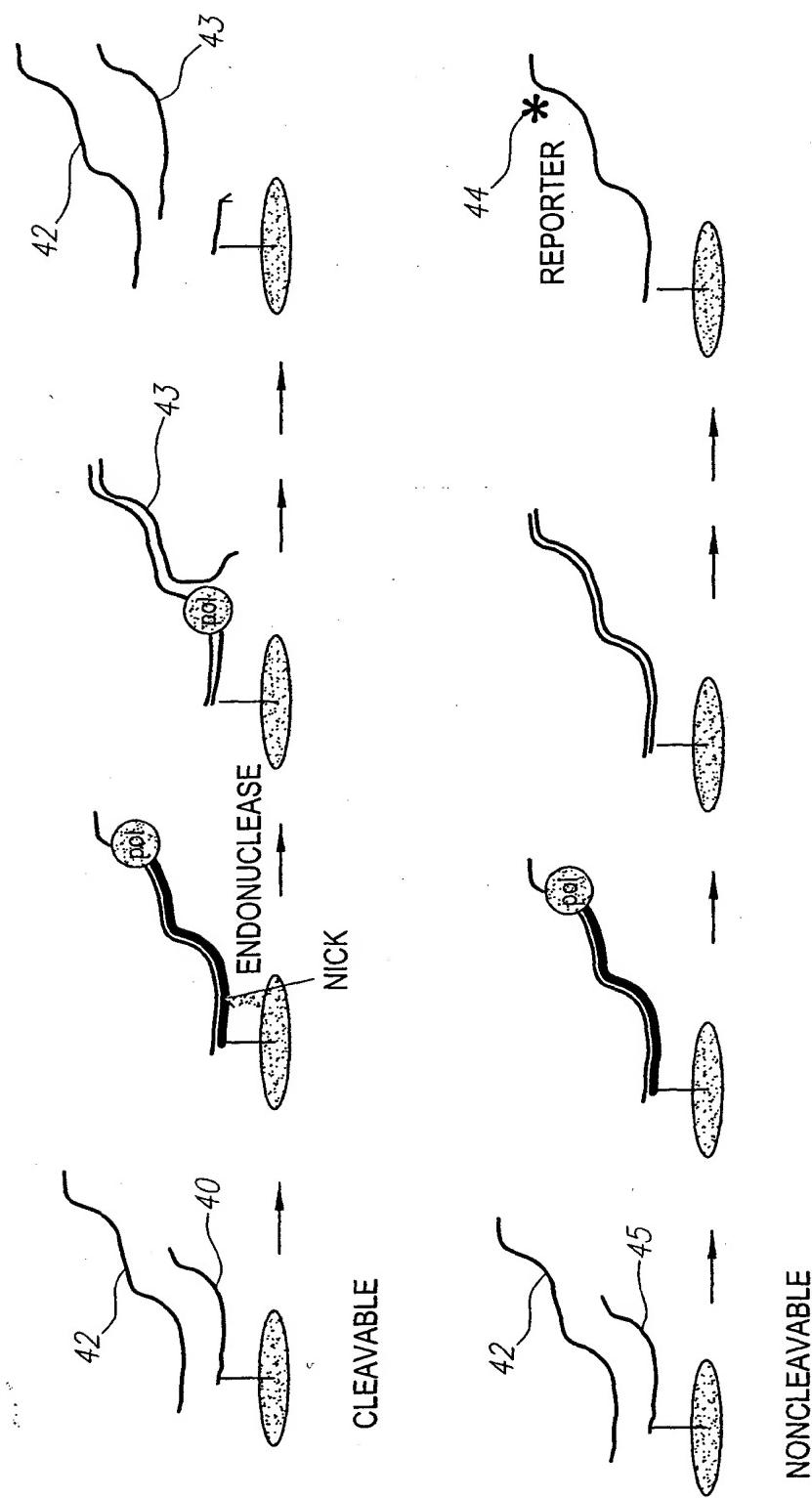
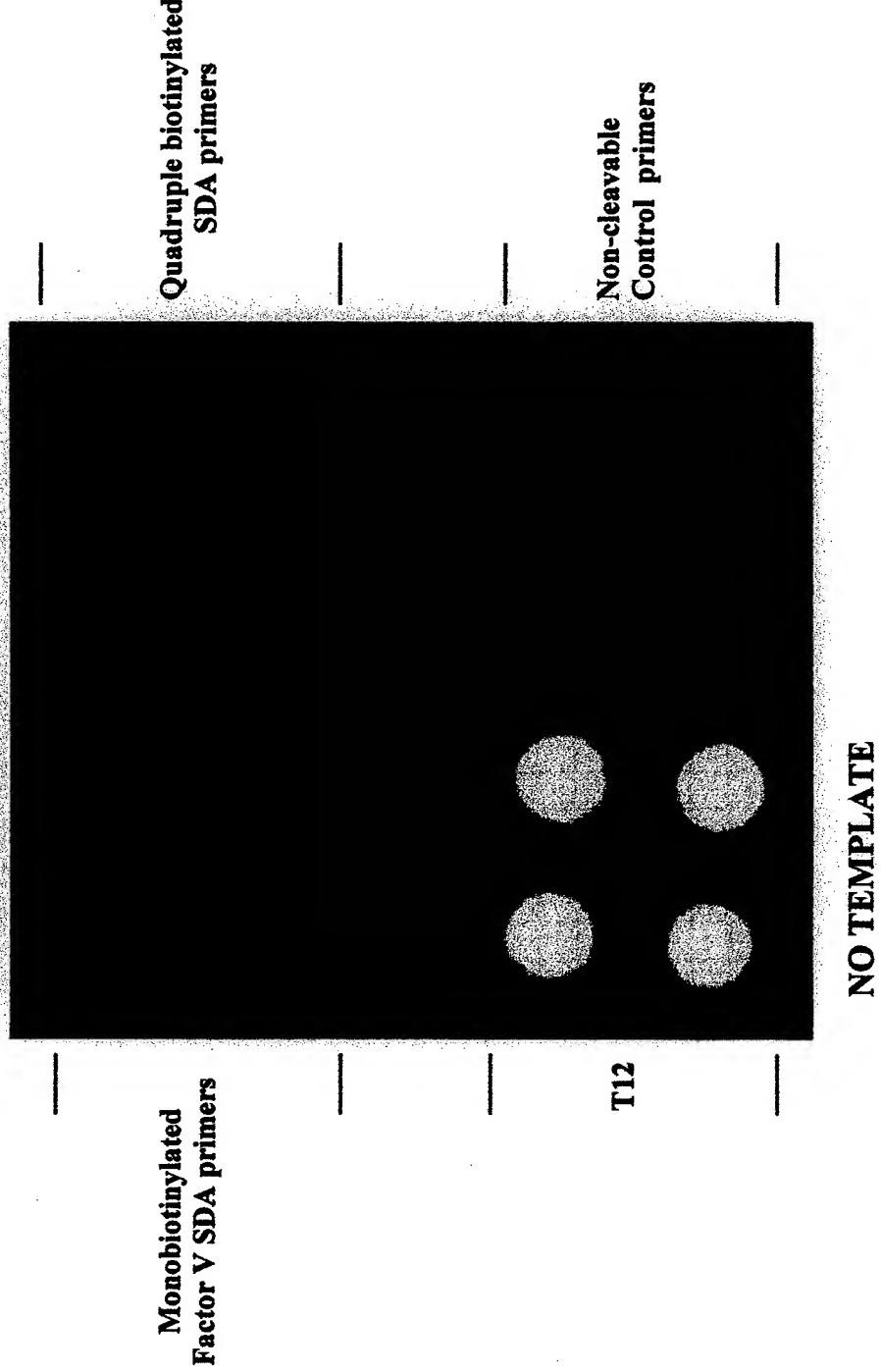


FIG. 5B

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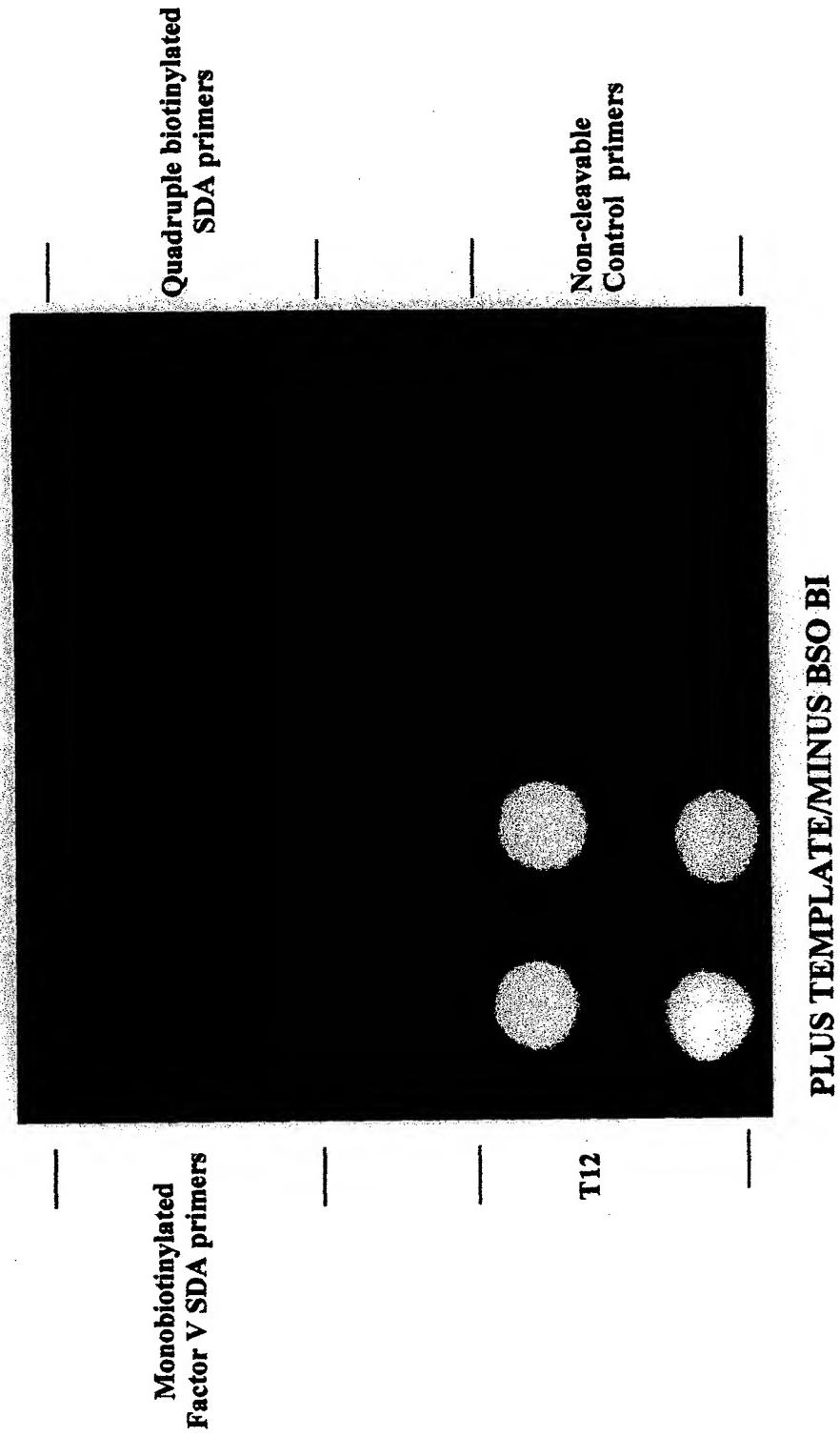
FIGURE 6A



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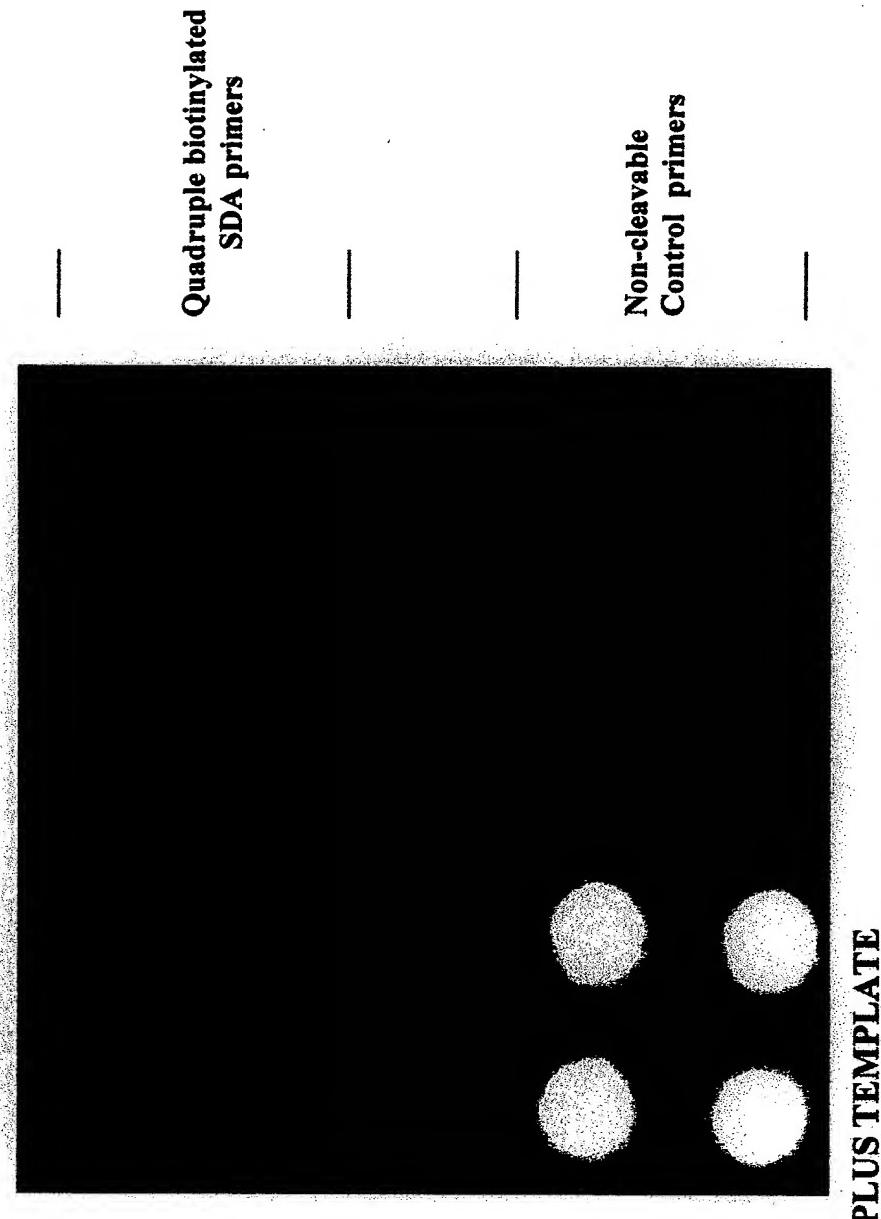
FIGURE 6B



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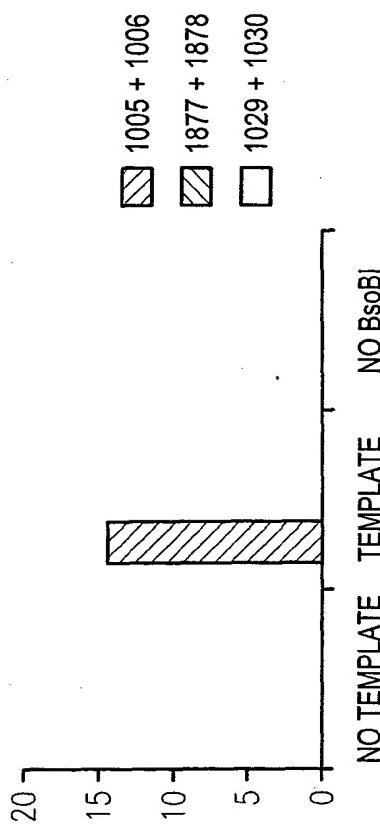
FIGURE 6C



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HUMAN COAGULATION FACTOR V ANCHORED SDA IN SITU
ON MICROCHIPS



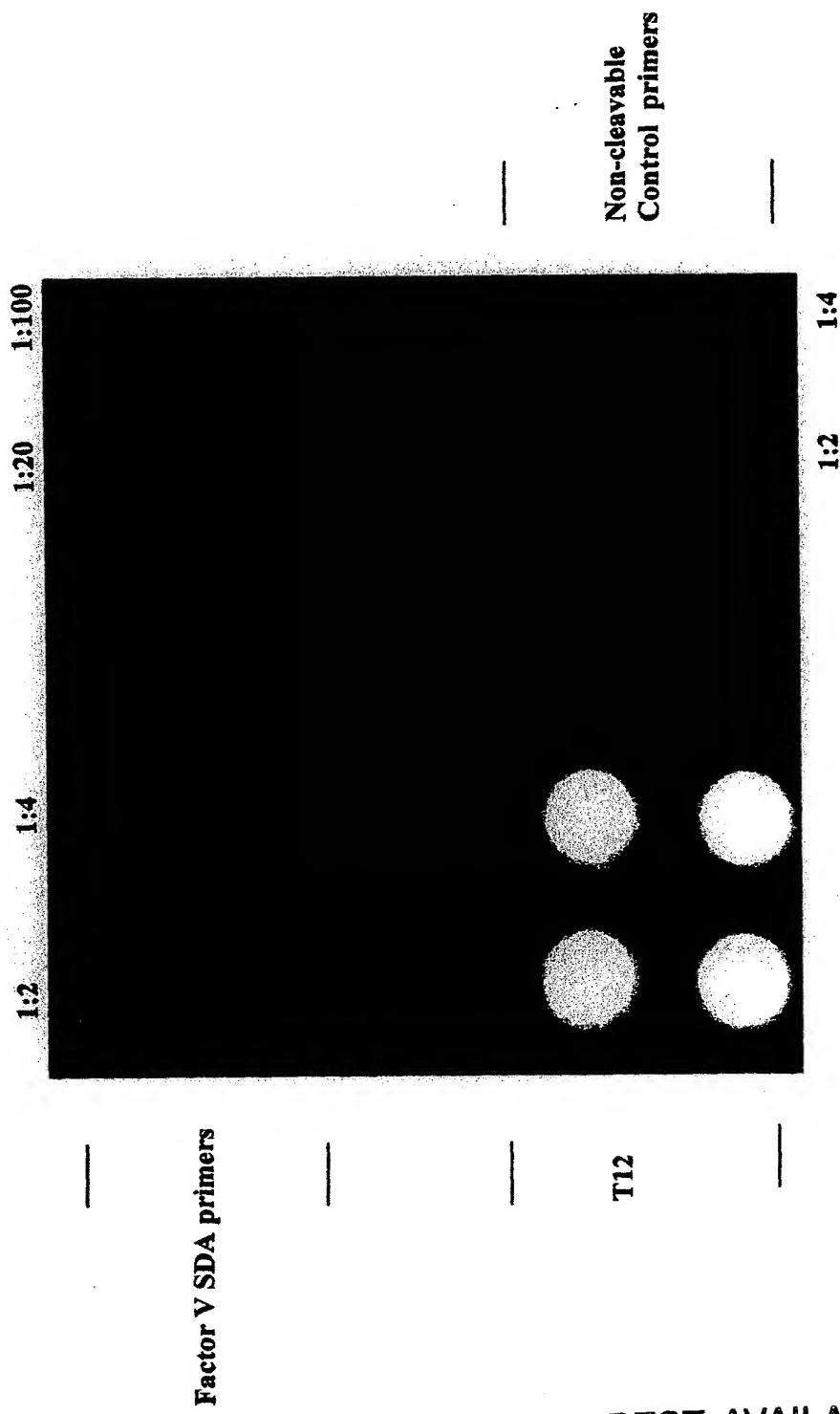
1005 + 1006 = MONOBIOTINYLATED PROBES
1877 + 1878 = QUADRUPLE-BIOTINYLATED PROBES
1029 + 1030 = NON-CLEAVABLE CONTROL PRIMERS

FIG. 7

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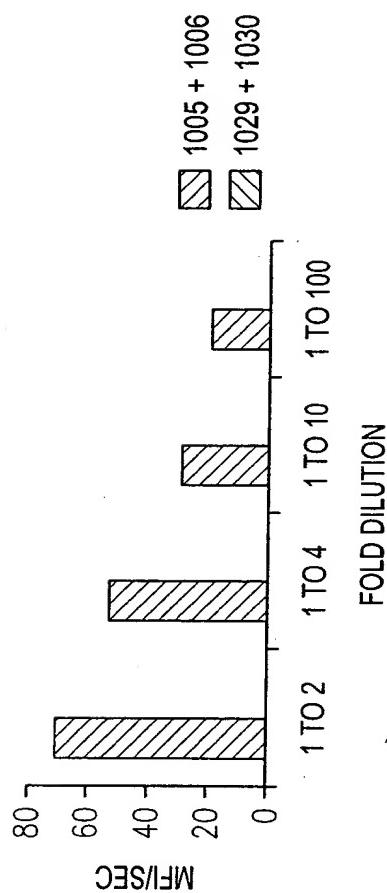
FIGURE 8



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TITRATION OF FACTOR V TEMPLATE IN
ANCHORED SDA



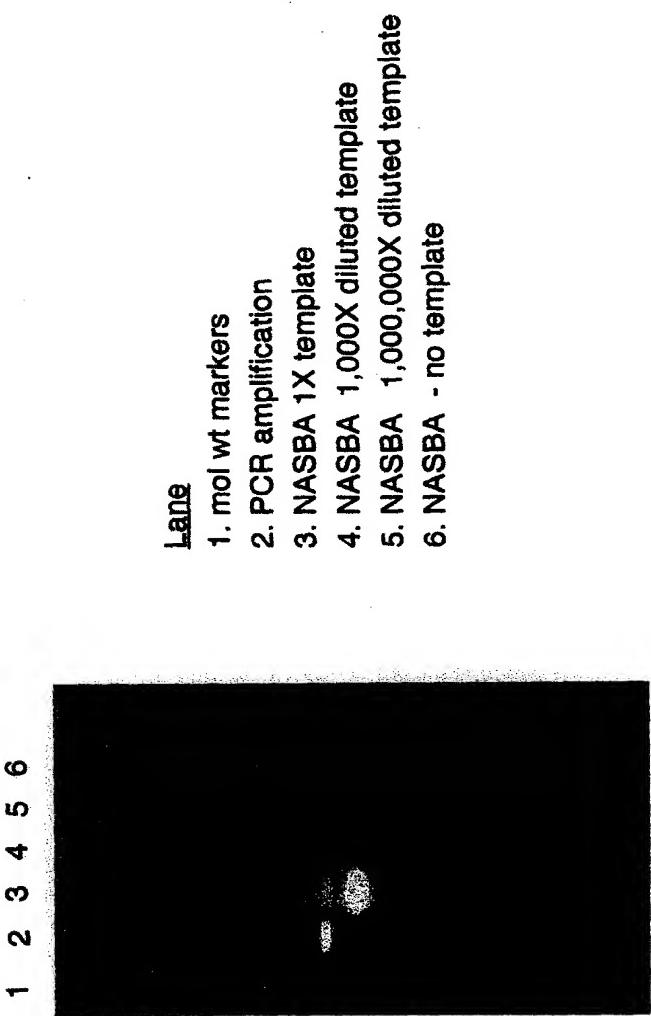
1005 + 1006 = FACTOR V SDA PRIMERS
1029 + 1030 = NON-CLEAVABLE CONTROL PRIMERS

FIG. 9

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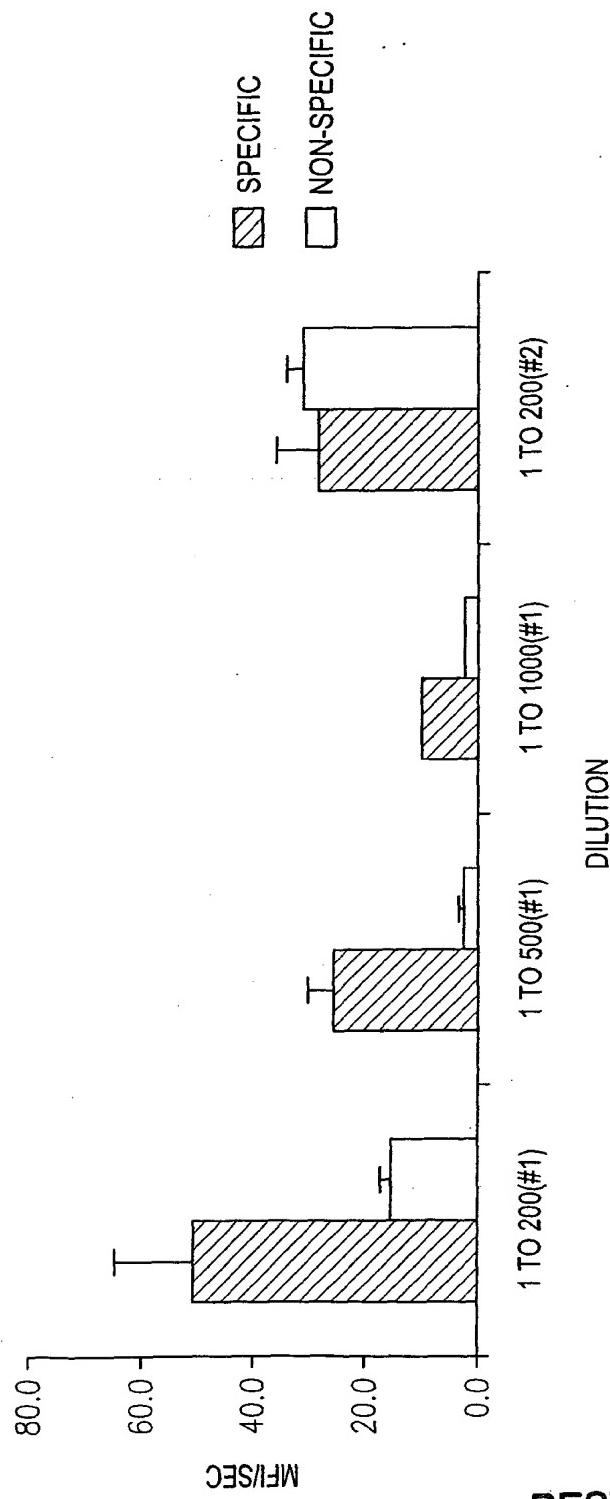
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FIGURE 10A



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FIG. 10B

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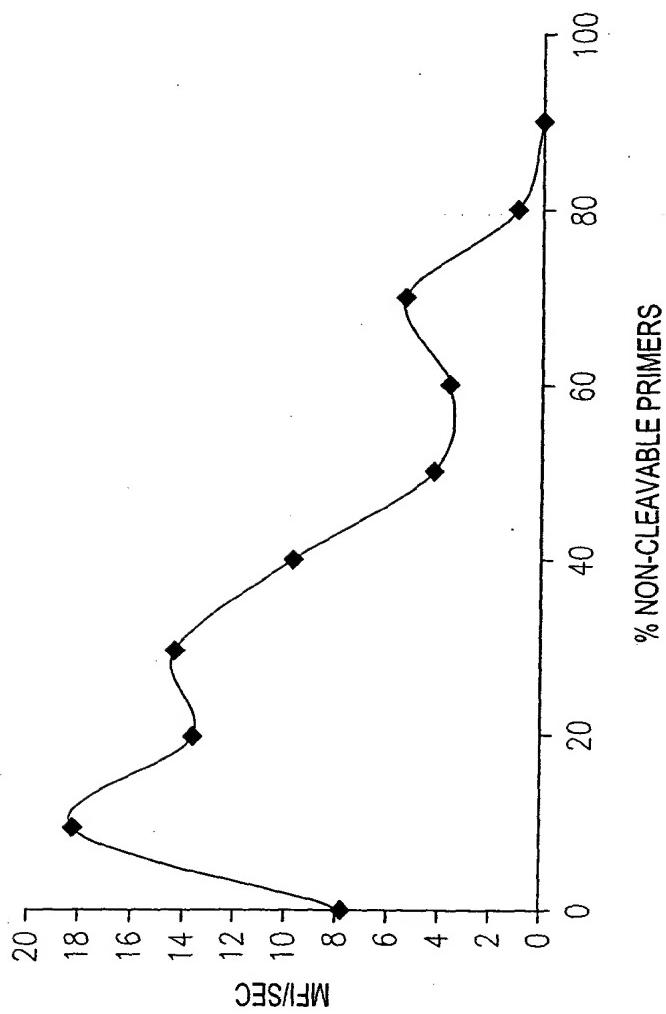


FIG. 11

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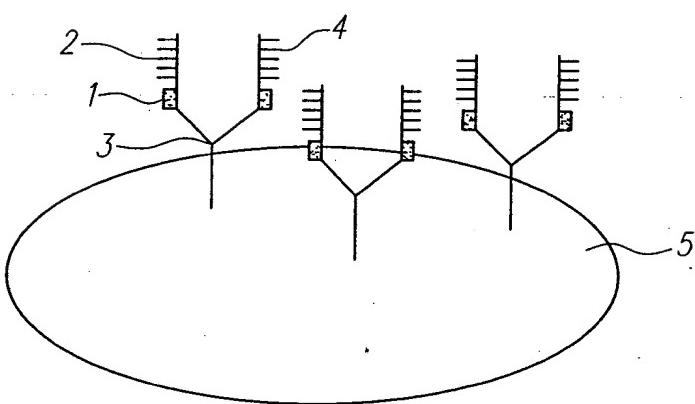


FIG. 12

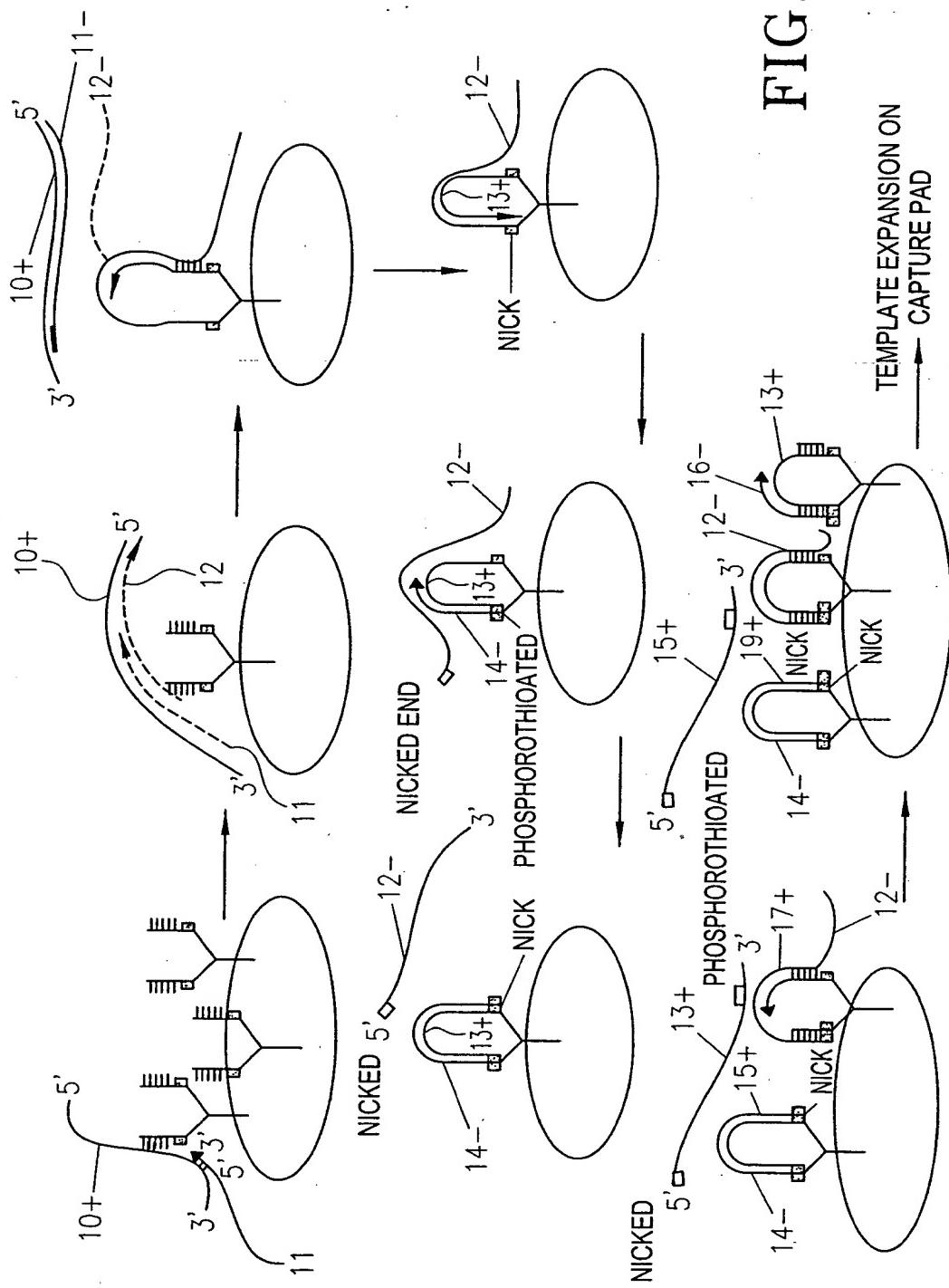
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09274685 "100901"

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FIG. 13

TEMPLATE EXPANSION ON
CAPTURE PAD



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09074168 - 1005001

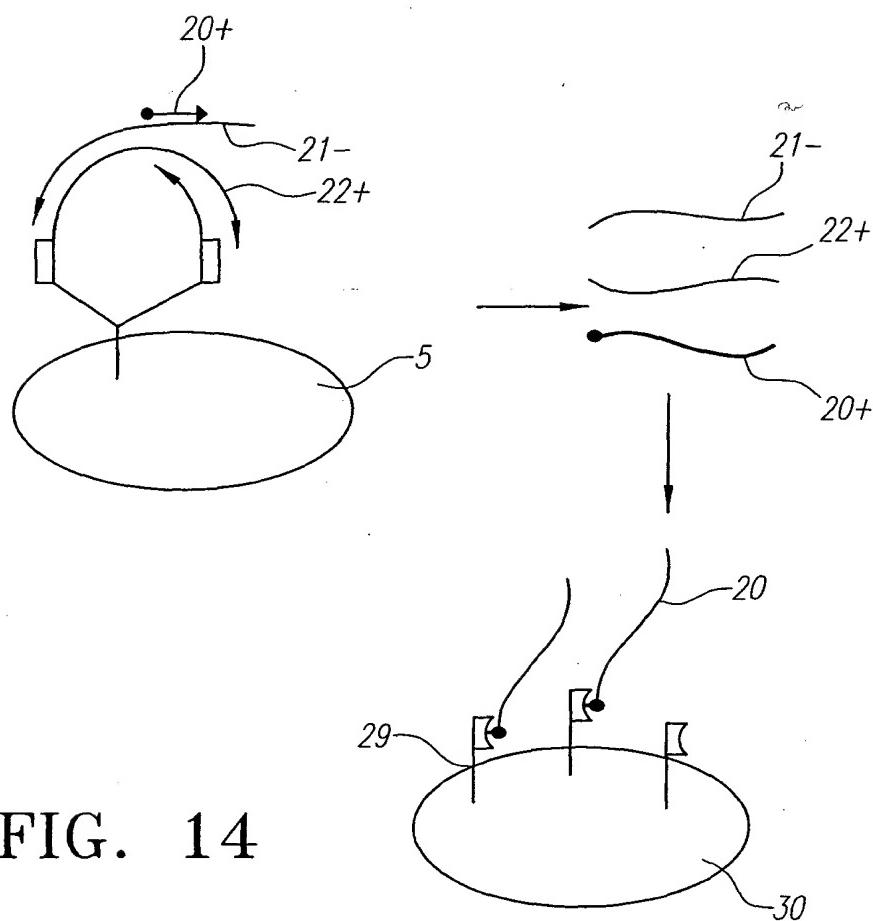


FIG. 14

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TOP SECRET - SECURITY INFORMATION

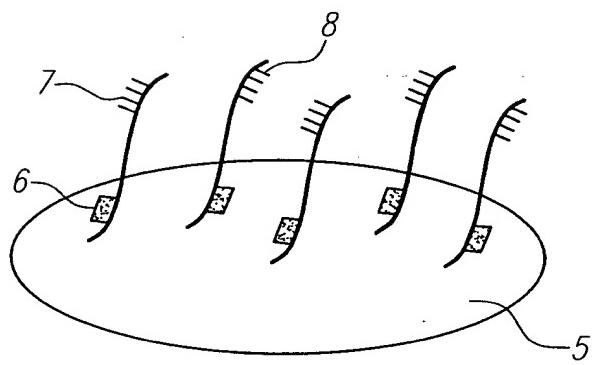
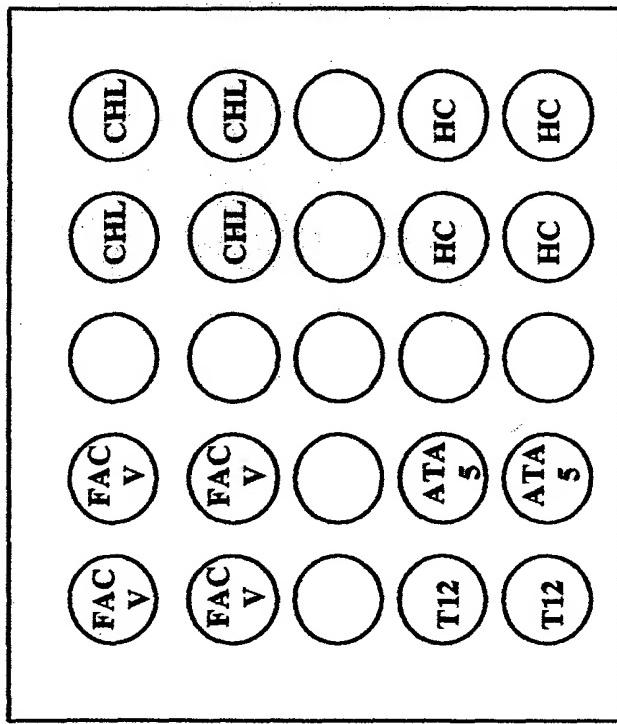


FIG. 15

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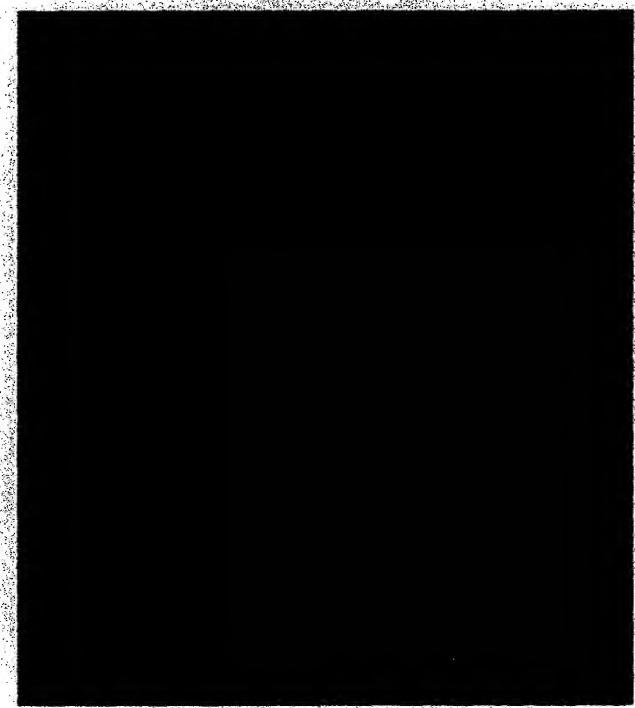
FIGURE 16



Experimental Layout

Control - No template +
all reporter oligos

FIGURE 17



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FIGURE 18

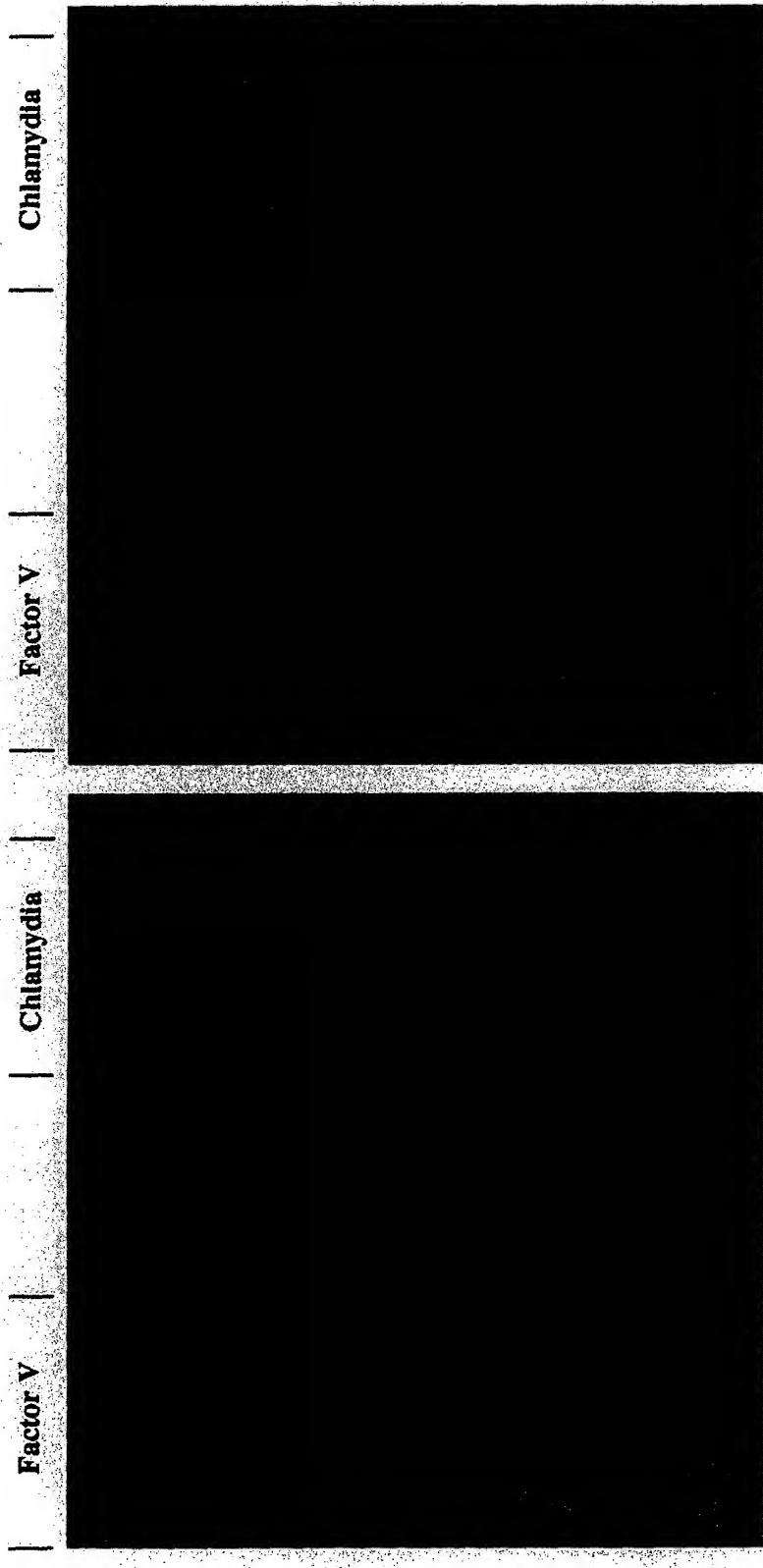


FIGURE 19



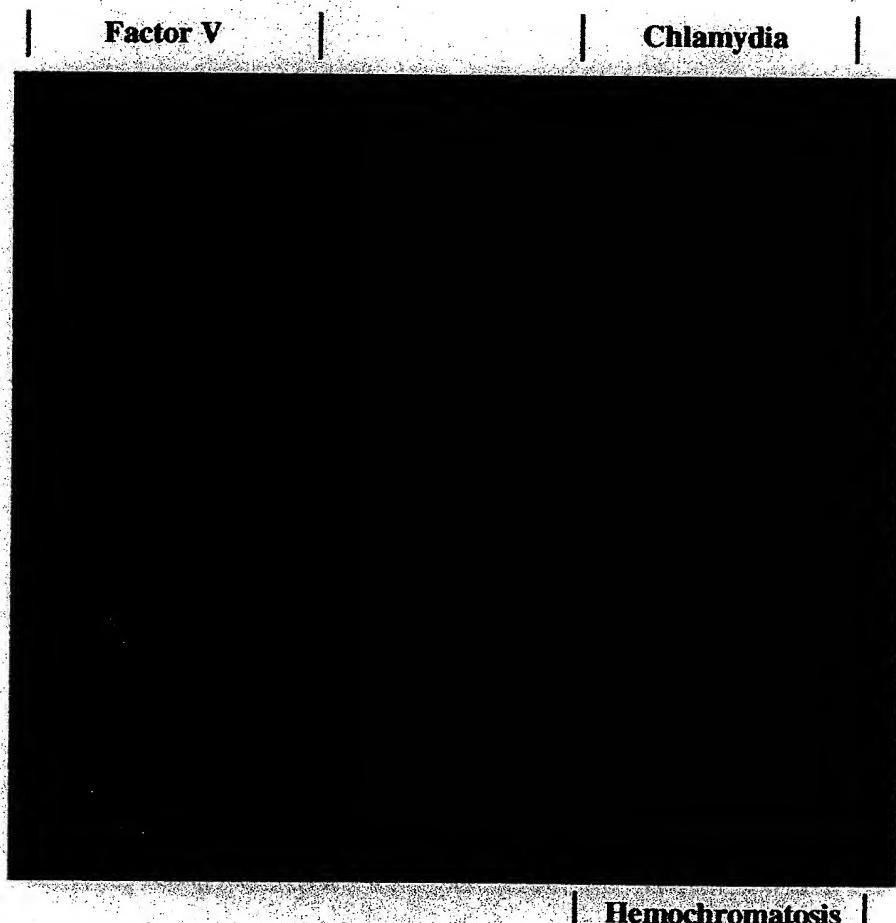
All templates + Factor V Reporter oligo

All templates + Factor V, Chlamydia
Reporter Oligos

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FIGURE 20

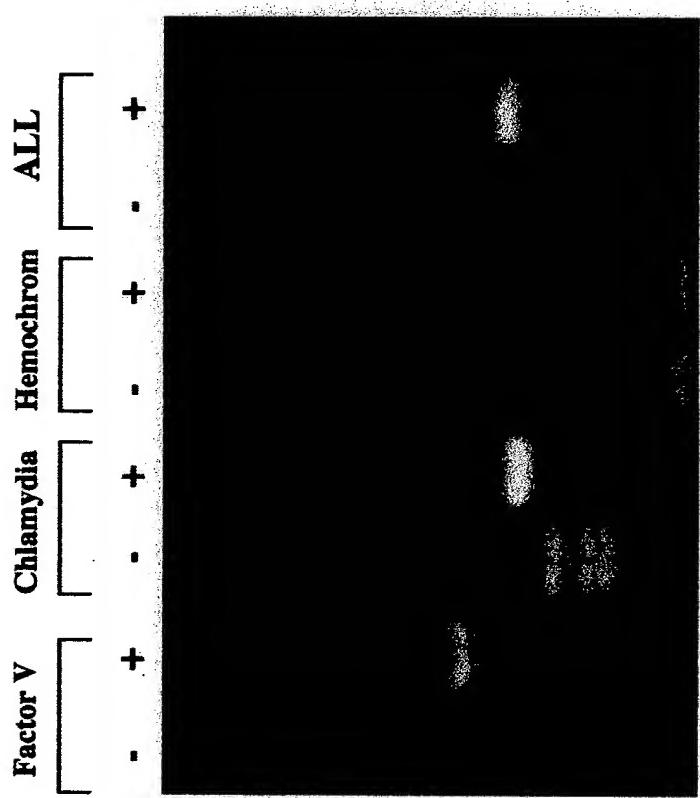


All templates + Factor V, Chlamydia and Hemachromatosis reporter oligos

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FIGURE 21

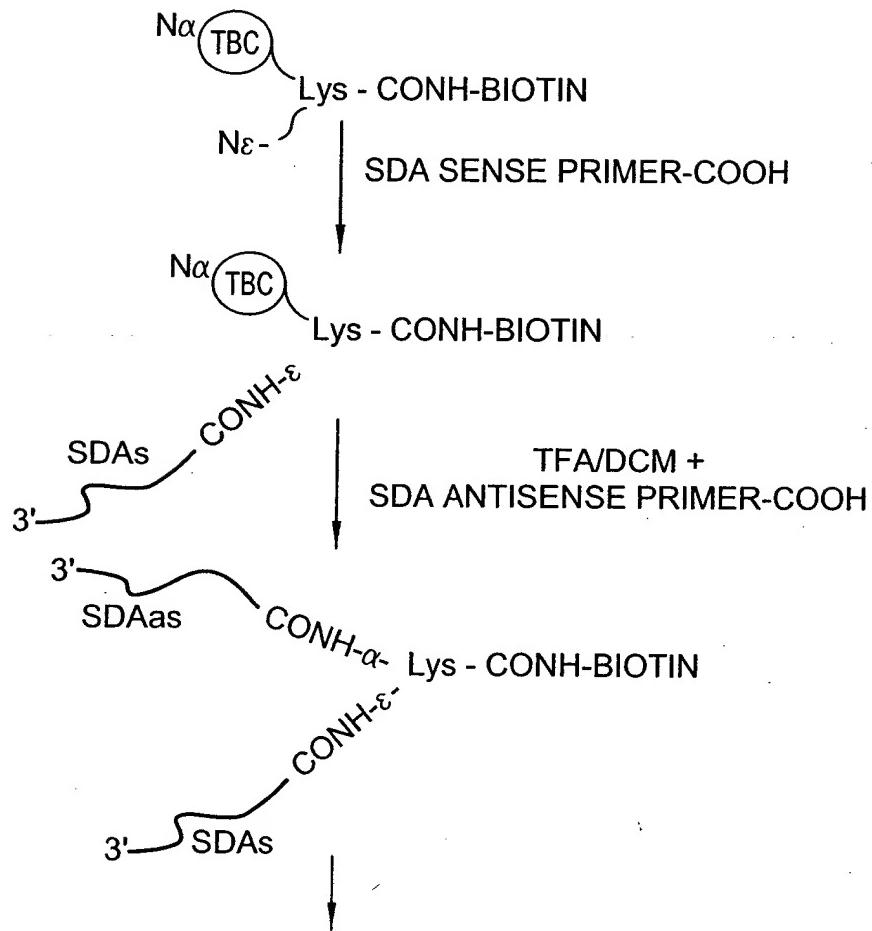


Control Solution SDA reactions

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FD600753942650



ATTACH TO STREPTAVIDIN PERMEATION LAYER ON MICROCHIP

FIG. 22

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SEQUENCE TO FOLLOW

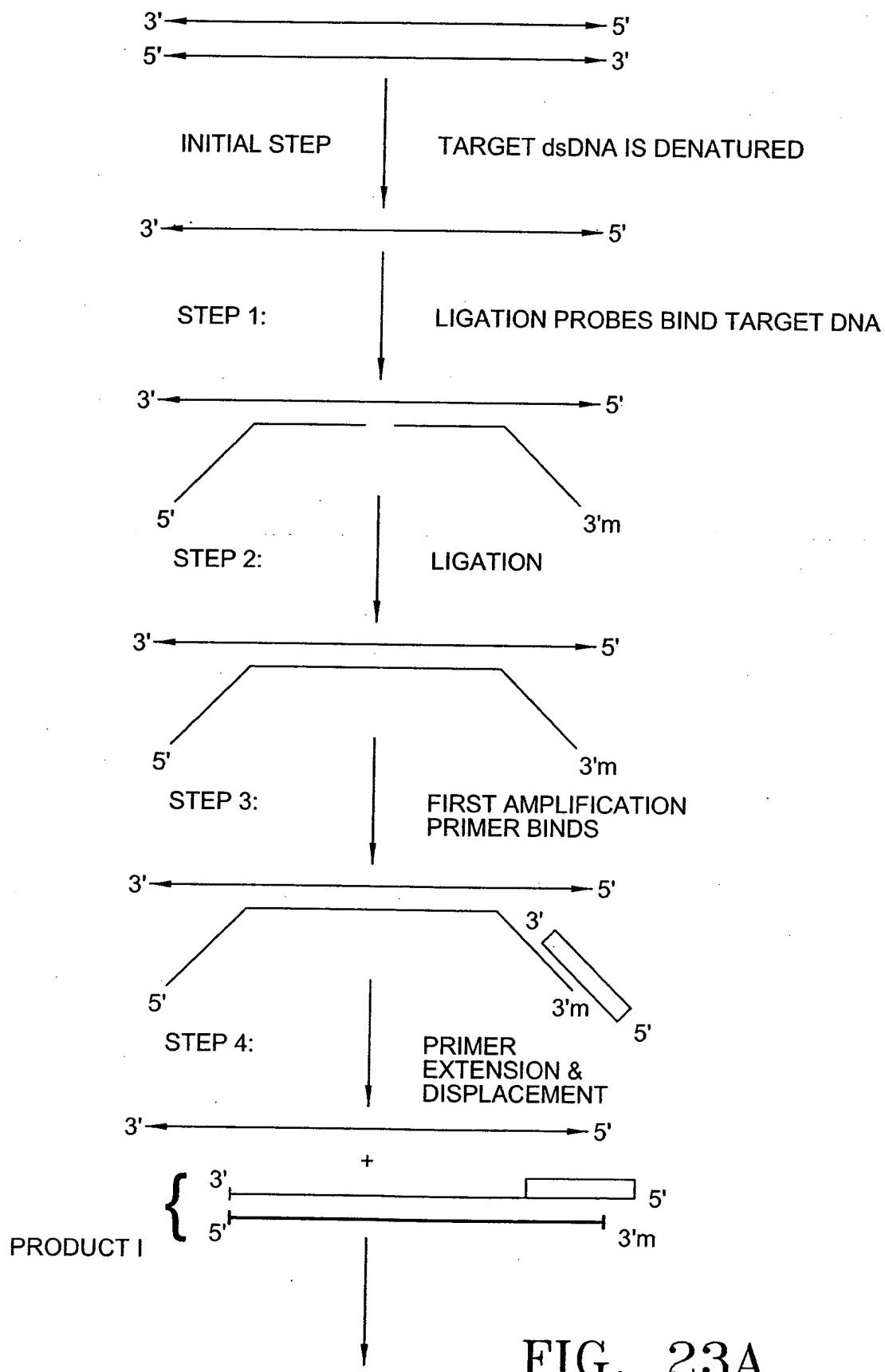


FIG. 23A

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TO6000T-580112660

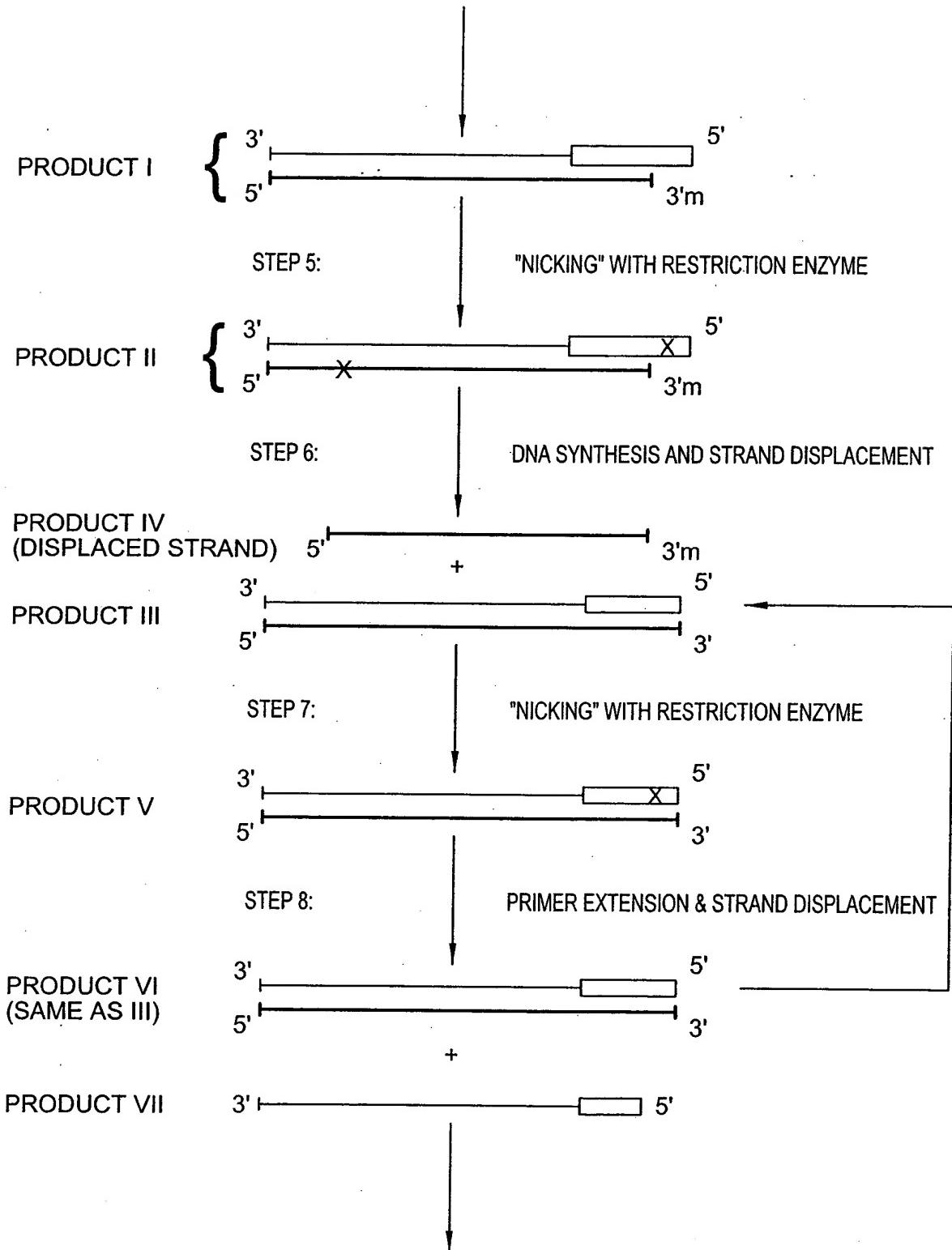
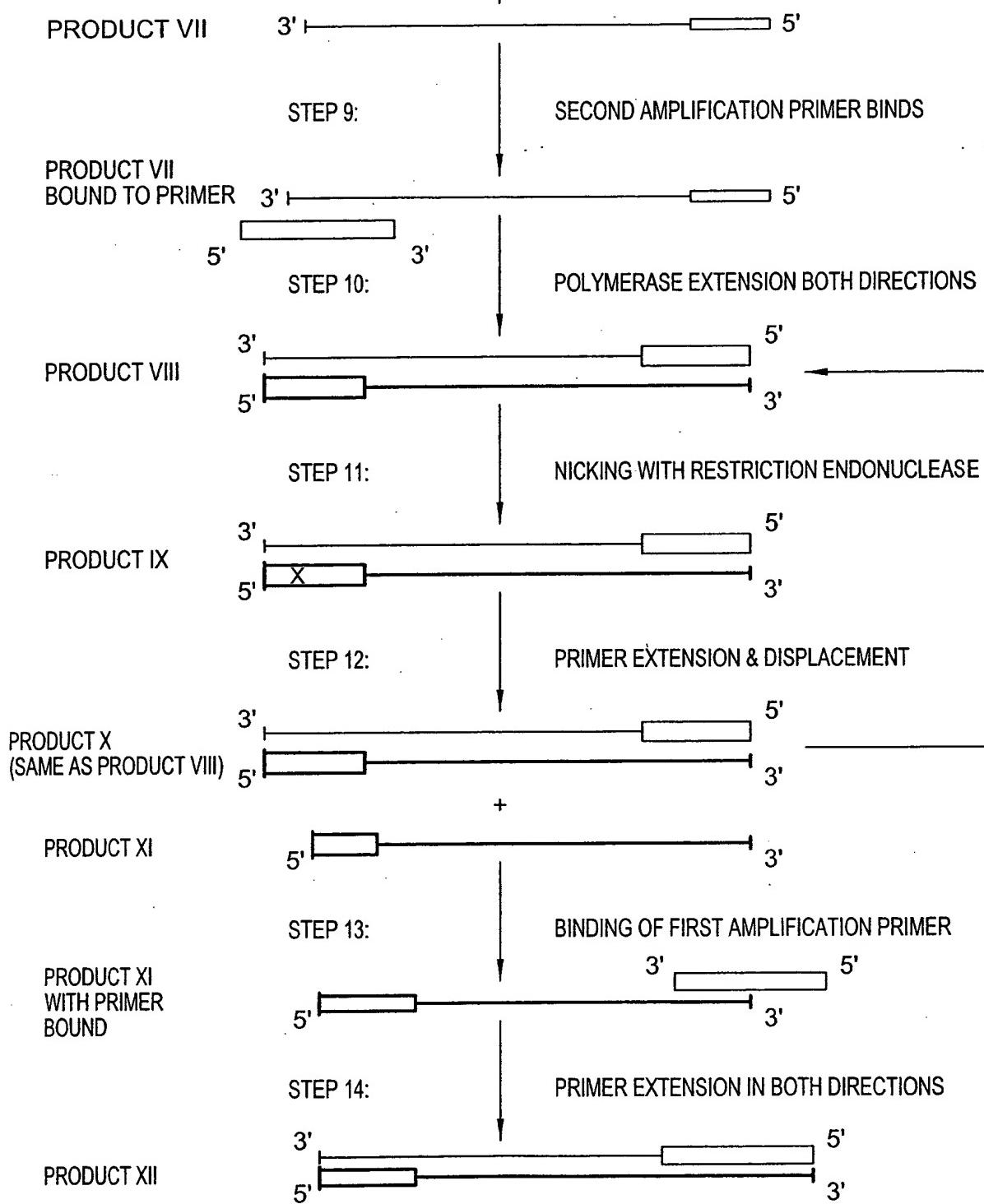


FIG. 23B

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09547488 - 106307



(PRODUCT XII CAN RE-ENTER PATHWAY AND BE FURTHER AMPLIFIED IN A MANNER
SIMILAR TO PRODUCT III, FOLLOWING STEP 6)

FIG. 23C BEST AVAILABLE COPY

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LIGATION-DEPENDENT DETECTION OF THE SALMONELLA spaQ GENE

GATION PROBES LP1 AND LP2:

spa ¹	TEMPLATE	5'-nnnnncaacatgacatcattacgagacggataatggatgttttagtgggg-3'
IPI ²		3'-*aattcccgcatgaggctggtaatgttgtactgttagtaatgttgtgc*-5'
		3'-ccatcaatttacctaataatcacgattatcccttagaagtcatgtgggctc ttcagacacctcgcctttagc-5' LP2 ³

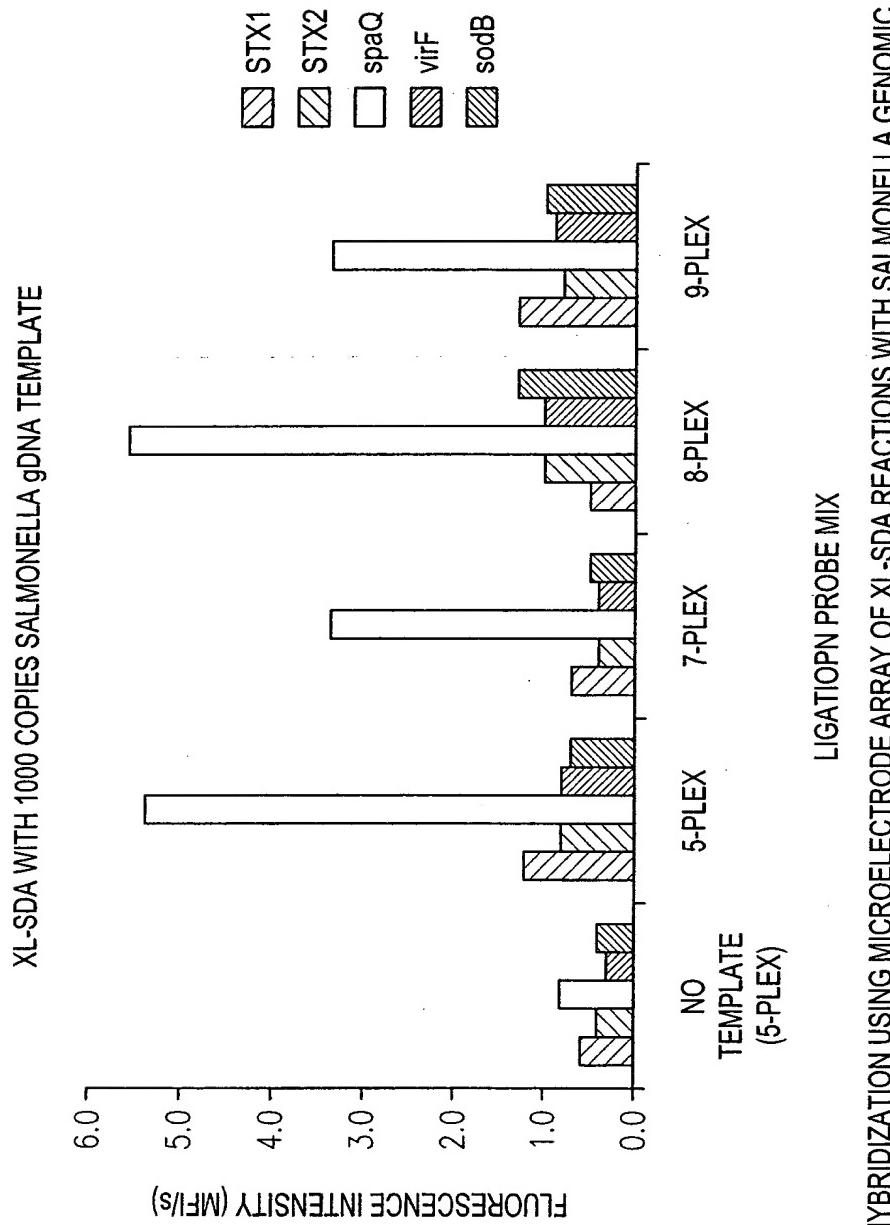
MPLIFICATION PRIMER SEQUENCES S1 AND S2:

<p>LPI 3'-<i>aattccgcattggtaatgttgactgttagtaatgctctgc</i>*-5'</p> <p style="text-align: center;"> </p> <p style="text-align: center;">5'-<i>accgcattttacctaataccaggattatcccctagagtcatgtggctttcagaccccgcccttagc</i>-5'</p>	<p>S1⁴</p>
<p>LP2 3'-<i>cctatcaattttacctaataccaggattatcccctagagtcatgtggctttcagaccccgcccttagc</i>-5'</p>	<p style="text-align: center;"> </p> <p style="text-align: center;">S2⁵</p>

FIG. 23D

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T D S O D T * 3 8 3 4 2 6 6 0



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FIG. 24

ELECTRONIC HYBRIDIZATION USING MICROELECTRODE ARRAY OF XL-SDA REACTIONS WITH SALMONELLA GENOMIC DNA AT 1000